Repairer Preventive Maintenance Checks and Services

6530-01-327-0686 Ventilator, Volume, Portable, Model 750M

[M-Monthly, Q-Quarterly, S-Semiannually, and A-Annually]

ITEM NO	INTERVAL	ITEM TO BE INSPECTED AND PROCEDURE	IS NOT MISSION CAPABLE IF:
1	S	Ventilator a. Conduct an inventory to ensure that the items listed on the Equipment Parts and Accessories List are on hand.	Missing components or accessories prevent the operation of the ventilator.
		b. Inspect hoses, fittings, and regulators for cracks, crimps, leakage, discoloration, damaged connector fittings, or general wear as directed in the manufacturer's literature.	Unserviceable accessories prevent use of the ventilator.
2	S	Preventative Maintenance Inspections a. Perform visual checks as directed in the manufacturer's literature.	The inspections do not pass standards.
		b. Perform performance checks as directed in the manufacturer's literature.	The inspections do not pass standards.
		c. Clean unit as directed in the manufacturer's literature.	
3	S	Case	
		Check for wear, loose or missing hardware, and cracks as directed in the manufacturer's literature.	The unserviceable case prevents protective storage, safe movement, or operation of the unit.
4	S	Battery a. Test the control module for operation using the internal battery as directed in the manufacturer's literature.	Use of the battery causes an alarm condition.
		 b. Check for a battery alarm as directed in the manufacturer's literature. 	
5	S	Multivoltage Power Supply a. Check the power supply for worn, cracked, or damaged connectors as directed in the manufacturer's literature.	The ventilator cannot be operated or if an electrical hazard is present.
		b. Test the operation of the power supply and the integrated battery charger as directed in the manufacturer's literature.	The multivoltage power supply is inoperable.
		c. Verify that electrical safety tests have been performed as scheduled as directed in the manufacturer's literature.	The unit fails any safety test.

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6	S	Patient Valve	
		Check for cracks, leakage, discoloration, and general wear as directed in the manufacturer's literature.	The patient valve is inoperable, malfunctioning, or endangers the patient.
7	S	Control Module	
		a. Check for tactile feel of all controls. Verify operation of controls as directed in the manufacturer's literature.	Any control is inoperable.
		b. Verify completion of self-test as directed in the manufacturer's literature.	Any portion of the self-test fails or aborts.
		c. Verify transducer calibration as directed in the manufacturer's literature.	The transducer fails calibration test.
		d. Check the various modes of operation as directed in the manufacturer's literature.	The ventilator does not operate in any of the modes of operation.
		(1) Verify the control ventilation – with/without "SIGH" - with/without "PEEP" as directed in the manufacturer's literature.	
		(2) Verify the assist-control ventilation – with/without "SIGH" – with/without "PEEP" as directed in the manufacturer's literature.	
		(3) Verify the synchronized intermittent mandatory ventilation (SIMV) – with/without "SIGH" – with/without "PEEP" as directed in the manufacturer's literature.	
		(4) Verify the assist-control backup during apnea – with/without "SIGH" – with/without "PEEP" as directed in the manufacturer's literature.	
		e. Update the Medical Equipment Verification/Certification sticker (DD Form 2163).	The unit has not been verified within the last six (6) months.